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=> s lysine(w)rich and seed and protein and zein
L1 4 LYSINE(W) RICH AND SEED AND PROTEIN AND ZEIN

=> d l1 1-4 ibib ab

L1 ANSWER 1 OF 4 AGRICOLA Compiled and distributed by the National
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(2004) on STN

ACCESSION NUMBER: 1998:49315 AGRICOLA

DOCUMENT NUMBER: IND21243126

TITLE: ***Lysine*** - ***rich*** modified gamma-
zeins accumulate in ***protein*** bodies
of transiently transformed maize endosperms.

AUTHOR(S): Torrent, M.; Alvarez, I.; Geli, M.I.; Dalcol, I.;
Ludevid, D.

AVAILABILITY: DNAL (QK710.P62)

SOURCE: Plant molecular biology, May 1997. Vol. 34, No. 1. p.
139-149

Publisher: Dordrecht : Kluwer Academic Publishers.

CODEN: PMBIDB; ISSN: 0167-4412

NOTE: Includes references

PUB. COUNTRY: Netherlands

DOCUMENT TYPE: Article

FILE SEGMENT: Non-U.S. Imprint other than FAO

LANGUAGE: English

AB During maize ***seed*** development, endosperm cells synthesize large
amounts of storage ***proteins***, alpha-, beta-, and gamma-
zeins, which accumulate within endoplasmic reticulum (ER)-derived
protein bodies. The absence of lysine in all ***zein***
polypeptides results in an imbalance in the amino acid composition of
maize ***seeds***. We modified the maize gamma- ***zein*** gene
through the introduction of ***lysine*** - ***rich*** (Pro-Lys)n
coding sequences at different sites of the gamma- ***zein*** coding
sequence. Maize endosperms were transiently transformed by biolistic
bombardment with Lys-rich gamma- ***zein*** constructs under the

Pro-Xaa region of the gamma-***zein***, high levels of ***protein*** were observed. In contrast, when (Pro-Lys)_n sequences were inserted five residues from the C-terminal, the transcript was present but modified ***protein*** was not detected. These results suggest that only an appropriate positioning of Lys-rich inserts leads to the modified molecule displaying correct folding and stability. Subcellular localization analyses and immunoelectron microscopy studies on isolated ***protein*** bodies demonstrated that modified gamma-***zeins*** accumulate within these organelles and co-localized with endogenous alpha- and gamma-***zeins***. The studies reported here show the feasibility of manipulating the gamma-***zein*** gene in order to obtain stable and correctly targeted Lys-rich ***zeins*** in maize ***seeds***.

L1 ANSWER 2 OF 4 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

ACCESSION NUMBER: 1997:320606 BIOSIS

DOCUMENT NUMBER: PREV199799611094

TITLE: ***Lysine*** - ***rich*** modified gamma-***zeins*** accumulate in ***protein*** bodies of transiently transformed maize endosperms.

AUTHOR(S): Torrent, Margarita; Alvarez, Inaki; Geli, M. Isabel; Dalcol, Ionara; Ludevid, Dolors [Reprint author]

CORPORATE SOURCE: Dep. de Genetia Molecular, Centre d'Investigacio i Desenvolupament, 08034 Barcelona, Spain

SOURCE: Plant Molecular Biology, (1997) Vol. 34, No. 1, pp. 139-149.

CODEN: PMBIDB. ISSN: 0167-4412.

DOCUMENT TYPE: Article

LANGUAGE: English

ENTRY DATE: Entered STN: 26 Jul 1997

Last Updated on STN: 26 Jul 1997

AB During maize ***seed*** development, endosperm cells synthesize large amounts of storage ***proteins***, alpha-, beta-, and gamma-***zeins***, which accumulate within endoplasmic reticulum (ER)-derived ***protein*** bodies. The absence of lysine in all ***zein*** polypeptides results in an imbalance in the amino acid composition of maize ***seeds***. We modified the maize gamma-***zein*** gene through the introduction of ***lysine*** - ***rich*** (Pro-Lys), coding sequences at different sites of the gamma-***zein*** coding sequence. Maize endosperms were transiently transformed by biolistic bombardment with Lys-rich gamma-***zein*** constructs under the control of the 1.7 kb gamma-***zein*** ***seed***-specific promoter and the cauliflower mosaic virus (CaMV) 35S promoter. When (Pro-Lys)_n sequences were inserted contiguous to or in substitution of the Pro-Xaa region of the gamma-***zein***, high levels of ***protein*** were observed. In contrast, when (Pro-Lys)_n sequences were inserted five residues from the C-terminal, the transcript was present but modified ***protein*** was not detected. These results suggest that only an appropriate positioning of Lys-rich inserts leads to the modified molecule displaying correct folding and stability. Subcellular localization analyses and immunoelectron microscopy studies on isolated ***protein*** bodies demonstrated that modified gamma-***zeins*** accumulate within these organelles and co-localized with endogenous alpha- and gamma-***zeins***. The studies reported here show the feasibility of manipulating the gamma-***zein*** gene in order to obtain stable and correctly targeted Lys-rich ***zeins*** in maize ***seeds***.

ACCESSION NUMBER: 1997:367431 CAPLUS
 DOCUMENT NUMBER: 127:92732
 TITLE: ***Lysine*** - ***rich*** modified .gamma.-
 zeins accumulate in ***protein*** bodies
 of transiently transformed maize endosperms
 AUTHOR(S): Torrent, Margarita; Alvarez, Inaki; Geli, M. Isabel;
 Dalcol, Ionara; Ludevid, Dolores
 CORPORATE SOURCE: Departament de Genetica Molecular, Centre
 d'Investigacio i Desenvolupament, (CSIC), Barcelona,
 08034, Spain
 SOURCE: Plant Molecular Biology (1997), 34(1), 139-149
 CODEN: PMBIDB; ISSN: 0167-4412
 PUBLISHER: Kluwer
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 AB During maize ***seed*** development, endosperm cells synthesize large
 amts. of storage ***proteins***, .alpha.-, .beta.-, and .gamma.-
 zeins, which accumulate within endoplasmic reticulum (ER)-derived
 protein bodies. The absence of lysine in all ***zein***
 polypeptides results in an imbalance in the amino acid compn. of maize
 seeds. We modified the maize .gamma.- ***zein*** gene through
 the introduction of ***lysine*** - ***rich*** (Pro-Lys)n coding
 sequences at different sites of the .gamma.- ***zein*** coding
 sequence. Maize endosperms were transiently transformed by biolistic
 bombardment with Lys-rich .gamma.- ***zein*** constructs under the
 control of the 1.7 kb .gamma.- ***zein*** ***seed***-specific
 promoter and the cauliflower mosaic virus (CaMV) 35S promoter. When
 (Pro-Lys)n sequences were inserted contiguous to or in substitution of the
 Pro-Xaa region of the .gamma.- ***zein***, high levels of
 protein were obsd. In contrast, when (Pro-Lys)n sequences were
 inserted five residues from the C-terminal, the transcript was present but
 modified ***protein*** was not detected. These results suggest that
 only an appropriate positioning of Lys-rich inserts leads to the modified
 mol. displaying correct folding and stability. Subcellular localization
 analyses and immunoelectron microscopy studies on isolated ***protein***
 bodies demonstrated that modified .gamma.- ***zeins*** accumulate
 within these organelles and co-localized with endogenous .alpha.- and
 .gamma.- ***zeins***. The studies reported here show the feasibility
 of manipulating the .gamma.- ***zein*** gene in order to obtain stable
 and correctly targeted Lys-rich ***zeins*** in maize ***seeds***.

L1 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1997:60561 CAPLUS
 DOCUMENT NUMBER: 126:87158
 TITLE: Characterization of the variability in lysine content
 for normal and opaque2 maize endosperm
 AUTHOR(S): Moro, Gloverson L.; Habben, Jeffrey E.; Hamaker, Bruce
 R.; Larkins, Brian A.
 CORPORATE SOURCE: Dep. Plant Sciences, Univ. Arizona, Tucson, AZ, 85721,
 USA
 SOURCE: Crop Science (1996), 36(6), 1651-1659
 CODEN: CRPSAY; ISSN: 0011-183X
 PUBLISHER: Crop Science Society of America, Inc.
 DOCUMENT TYPE: Journal

increasing the content of this essential amino acid in endosperm
 proteins depends on understanding the mechanisms regulating the
 synthesis and accumulation of ***lysine*** - ***rich***
 proteins. The variability for lysine and ***protein***
 contents was studied in maize endosperm. Amts. of total ***protein***
 , ***zeins*** , and non- ***zeins*** measured by microKjeldahl, and
 lysine content, estd. by amino acid anal., were detd. for 93 maize
 inbreds. Addnl., an ELISA was used to est. the relative content of the
 protein synthesis factor EF-1.alpha. in 20 selected genotypes.
 Considerable differences in lysine content were obsd. among normal and
 opaque2 genotypes, with the effect of the mutation being highly dependent
 on the genetic background. A high correlation was detected between the
 lysine content and the concn. of total non- ***zein*** ***proteins***
 and EF-1.alpha.. An assay for EF-1.alpha. concn. may provide a simple and
 inexpensive method from breeding programs to select for improved
 protein quality.

REFERENCE COUNT: 44 THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s seed and protein and gamma and zein

L2 91 SEED AND PROTEIN AND GAMMA AND ZEIN

=> duplicate remove l2

DUPLICATE PREFERENCE IS 'AGRICOLA, BIOSIS, EMBASE, CAPLUS'

KEEP DUPLICATES FROM MORE THAN ONE FILE? Y/(N):n

PROCESSING COMPLETED FOR L2

L3 59 DUPLICATE REMOVE L2 (32 DUPLICATES REMOVED)

=> d l3 1-10

L3 ANSWER 1 OF 59 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2004:174715 CAPLUS

TI Matrix-Assisted Laser Desorption Ionization Time-of-Flight Mass
 Spectrometry Analysis of ***Zeins*** in Mature Maize Kernels

AU Adams, Whitney R.; Huang, Shihshieh; Kriz, Alan L.; Luethy, Michael H.

CS Mystic Research, Monsanto Company, Mystic, CT, 06355, USA

SO Journal of Agricultural and Food Chemistry (2004), 52(7), 1842-1849

CODEN: JAFCAU; ISSN: 0021-8561

PB American Chemical Society

DT Journal

LA English

RE.CNT 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 59 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2003:242097 CAPLUS

DN 138:267201

TI Pesticidal compositions for coating plant propagation material containing
 anthranilamides

IN Berger, Richard Alan; Flexner, John Lindsey

PA E. I. Du Pont de Nemours & Co., USA

SO PCT Int. Appl., 147 pp.

CODEN: PIXXD2

=> d 13 36 ibib ab

L3 ANSWER 36 OF 59 AGRICOLA Compiled and distributed by the National
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(2004) on STN DUPLICATE 11

ACCESSION NUMBER: 97:33751 AGRICOLA
DOCUMENT NUMBER: IND20564663
TITLE: The maize ***gamma*** - ***zein*** sequesters
alpha- ***zein*** and stabilizes its accumulation
in ***protein*** bodies of transgenic tobacco
endosperm.
AUTHOR(S): Coleman, C.E.; Herman, E.M.; Takasaki, K.; Larkins,
B.A.
CORPORATE SOURCE: Brigham Young University, Provo, UT.
AVAILABILITY: DNAL (QK725.P532)
SOURCE: The Plant cell, Dec 1996. Vol. 8, No. 12. p. 2335-2345
Publisher: [Rockville, MD : American Society of Plant
Physiologists, c1989-
CODEN: PLCEEW; ISSN: 1040-4651
NOTE: Includes references
PUB. COUNTRY: Maryland; United States
DOCUMENT TYPE: Article
FILE SEGMENT: U.S. Imprints not USDA, Experiment or Extension
LANGUAGE: English

AB ***Zeins*** are ***seed*** storage ***proteins*** that form
accretions called ***protein*** bodies in the rough endoplasmic
reticulum of maize endosperm cells. Four types of ***zeins*** , alpha,
beta, ***gamma*** , and delta, aggregate in a distinctive spatial
pattern within the ***protein*** body. We created transgenic tobacco
plants expressing alpha- ***zein*** , ***gamma*** - ***zein*** , or
both to examine the interactions between these ***proteins*** leading
to the formation of ***protein*** bodies in the endosperm. Whereas
gamma - ***zein*** accumulated in ***seeds*** of these
plants, stable accumulation of alpha- ***zein*** required simultaneous
synthesis of ***gamma*** - ***zein*** . The ***zein***
proteins formed accretions in the endoplasmic reticulum similar
to
those in maize endosperm. ***Protein*** bodies were also found in
protein storage vacuoles. The accumulation of both types of
zeins peaked early in development and declined during maturation.
Even in the presence of ***gamma*** - ***zein*** , there was a
turnover of alpha- ***zein*** , suggesting that the interaction between
the two ***proteins*** might be transitory. We suggest that
gamma - ***zein*** plays an important role in ***protein***
body formation and demonstrate the utility of tobacco for studying
interactions between different ***zeins*** .

=> s gamma(w)zein and transform?

L4 35 GAMMA(W) ZEIN AND TRANSFORM?

FILES SUBMITTED FROM MORE THAN ONE FILE: 1/ (N) : 11
PROCESSING COMPLETED FOR L4

L5 20 DUPLICATE REMOVE L4 (15 DUPLICATES REMOVED)

=> d l5 1-20 ti

L5 ANSWER 1 OF 20 CAPLUS COPYRIGHT 2004 ACS on STN

TI Production of peptides and proteins by accumulation in plant endoplasmic reticulum-derived protein bodies

L5 ANSWER 2 OF 20 CAPLUS COPYRIGHT 2004 ACS on STN

TI Self-processing transgenic plants and plant parts expressing hyperthermophilic processing enzymes

L5 ANSWER 3 OF 20 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN DUPLICATE 1

TI Expression of the ***gamma*** - ***zein*** protein of maize in seeds of transgenic barley: effects on grain composition and properties.

L5 ANSWER 4 OF 20 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN DUPLICATE 2

TI Zein accumulation in forage species (Lotus corniculatus and Medicago sativa) and co-expression of the ***gamma*** - ***zein*** :KDEL and beta-zein: KDEL polypeptides in tobacco leaf.

L5 ANSWER 5 OF 20 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN DUPLICATE 3

TI Combination of viral promoter sequences to generate highly active promoters for heterologous therapeutic protein over-expression in plants.

L5 ANSWER 6 OF 20 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN

TI Expression of a synthetic E. coli heat-labile enterotoxin B sub-unit (LT-B) in maize.

L5 ANSWER 7 OF 20 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN DUPLICATE 4

TI Expression of a synthetic porcine alpha-lactalbumin gene in the kernels of transgenic maize.

L5 ANSWER 8 OF 20 CAPLUS COPYRIGHT 2004 ACS on STN

TI Geminivirus replicases and the genes encoding them and their use to create polyploid plant cells

L5 ANSWER 9 OF 20 CAPLUS COPYRIGHT 2004 ACS on STN

TI Methods of using viral replicase polynucleotides and polypeptides in transgenic plants

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Terms	Documents
L7 and lysine adj enriched	5

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 Derwent World Patents Index
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 result set

DB=USPT,USOC,EPAB,JPAB,DWPI; PLUR=YES; OP=OR

<u>L8</u>	L7 and lysine adj enriched	5	<u>L8</u>
<u>L7</u>	zein and lysine and enriched	97	<u>L7</u>
<u>L6</u>	maize and increased adj lysine	44	<u>L6</u>
<u>L5</u>	maize and increased adj lusine	0	<u>L5</u>

DB=DWPI; PLUR=YES; OP=OR

<u>L4</u>	L3 and proline	1	<u>L4</u>
<u>L3</u>	wo adj 9312230	5	<u>L3</u>
<u>L2</u>	WO adj 9315221	4	<u>L2</u>
<u>L1</u>	WO 9315221	915453	<u>L1</u>

END OF SEARCH HISTORY